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OXIDIZING COMPOSITION AND USES FOR DYEING, FOR
PERMANENTLY RESHAPING OR FOR BLEACHING KERATIN FIBRES

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The present invention relates to an oxidizing
5 composition intended for treating keratin fibres,
comprising at least one enzyme of 2-electron
oxidoreductase type in the presence of at least one
donor for the said enzyme and at least one
aminosilicone as well as to its uses for dyeing, for
10 permanently reshaping or for bleaching keratin fibres,
in particular human hair.

It is known to dye keratin fibres, and in
particular human hair, with dye compositions containing
oxidation dye precursors, in particular para-
15 phenylenediamines, ortho- or para-aminophenols and
heterocyclic bases which are generally referred to as
oxidation bases. Oxidation dye precursors, or oxidation
bases, are colourless or weakly coloured compounds
which, when combined with oxidizing products, can give
20 rise to coloured compounds and dyes by a process of
oxidative condensation.

It is also known that the shades obtained with
these oxidation bases can be varied by combining them
with couplers or colour modifiers, the latter being
25 chosen in particular from aromatic meta-diamines, meta-
aminophenols, meta-diphenols and certain heterocyclic
compounds.

The variety of compounds used as regards the
oxidation bases and the couplers allows a wide range of
30 colours to be obtained.

The so-called "permanent" coloration obtained
by means of these oxidation dyes must moreover satisfy
a certain number of requirements. Thus it must have no
toxicological drawbacks, it must be able to give shades
35 of the desired intensity and it must be able to
withstand external agents (light, bad weather, washing,
permanent-waving, perspiration, rubbing).

The dyes must also be able to cover white hair
and, lastly, they must be as unselective as possible,
40 i.e. they must give the smallest possible colour

differences along the same length of keratin fibre, which may in fact be differently sensitized (i.e. damaged) between its tip and its root.

The oxidation dyeing of keratin fibres is generally carried out in alkaline medium, in the presence of hydrogen peroxide. However, the use of alkaline media in the presence of hydrogen peroxide has the drawback of causing appreciable degradation of the fibres, as well as considerable bleaching of the keratin fibres, which is not always desirable.

The oxidation dyeing of keratin fibres can also be carried out using oxidizing systems other than hydrogen peroxide, such as enzymatic systems. Thus, it has already been proposed to dye keratin fibres, in particular in patent application EP-A-0,310,675, with compositions comprising an oxidation dye precursor in combination with enzymes such as pyranose oxidase, glucose oxidase or uricase, in the presence of a donor for the said enzymes. Although being used under conditions which do not result in degradation of the keratin fibres which is comparable to that caused by the dyes used in the presence of hydrogen peroxide, these dye formulations nevertheless lead to colorations which are still insufficient, both as regards the homogeneity of the colour distributed along the fibre ("unison") and as regards the chromaticity (luminosity), the dyeing power and the resistance to the various aggressive factors to which the hair may be subjected.

It is known that the most common technique for obtaining a permanent reshaping of the hair consists, in a first stage, in opening the keratin -S-S-disulphide (cysteine) bonds using a composition containing a suitable reducing agent (reduction step) followed, after having rinsed the hair thus treated, by reconstituting, in a second stage, the said disulphide bonds by applying to the hair, which has been placed under tension beforehand (curlers and the like), an oxidizing composition (oxidation step, also known as the fixing

The reducing compositions which may be used in order to carry out the first step of a permanent-waving operation generally contain, as reducing agents, sulphites, bisulphites, alkylphosphines or, preferably, thiols. Among the thiols, those commonly used are cysteine and the various derivatives thereof, cysteamine and the derivatives thereof, thiolactic acid or thioglycolic acid, the salts thereof and the esters thereof, in particular glyceryl thioglycolate.

The problem of the technique of the permanent-
waving operations known to date is that their
application to the hair induces long-term adverse
changes in the quality of the hair. The essential
causes of these adverse changes in the quality of the
hair are a reduction in its cosmetic properties, such
as its sheen and its feel, and degradation of its
mechanical properties, more particularly degradation of
its mechanical strength due to swelling of the keratin
fibres during the rinsing between the reduction step
and the oxidation step, which can also be reflected by
an increase in its porosity. The hair is weakened and
can become brittle during subsequent treatments such as
blow-drying.

The same problem of adverse changes in keratin fibres is encountered during processes for bleaching the hair.

It is known that the permanent reshaping or bleaching of keratin fibres can also be carried out under milder conditions using oxidizing systems other than hydrogen peroxide, such as enzymatic systems. Thus, processes for the permanent reshaping or bleaching of keratin fibres have already been proposed, in particular in patent application EP-A-0,310,675, with compositions comprising an enzyme such as pyranose oxidase, glucose oxidase or uricase, in the presence of a donor for the said enzyme. Although being used under conditions which do not result in degradation of the keratin fibres which is comparable to that caused by conventional permanent-waving or bleaching processes, these oxidizing formulations nevertheless lead to results which are still insufficient, as regards the curl hold over time, as regards the compatibility of permanent-waved or bleached hair with subsequent treatments, as regards the degradation of the mechanical properties of the permanent-waved hair, in particular the reduction of the porosity of the hair, and as regards the reduction of the cosmetic properties such as the feel, or alternatively as regards the uniformity of the bleaching along the keratin fibres.

The aim of the present invention is to solve the problems mentioned above.

The Applicant has discovered, surprisingly, novel compositions containing, as oxidizing system, at least one enzyme of 2-electron oxidoreductase type in the presence of at least one donor for the said enzyme and at least one aminosilicone, which can constitute, in the presence of oxidation dye precursors (oxidation bases) and optionally couplers, ready-to-use dye formulations which lead to more homogeneous, more intense and more chromatic colorations without giving rise to any significant degradation, these colorations being relatively unselective and showing good

The Applicant has also discovered, unexpectedly, that the use, in a process for the permanent reshaping of keratin fibres, of an oxidizing composition containing, as oxidizing system, at least one enzyme of 2-electron oxidoreductase type in the presence of at least one donor for the said enzyme and at least one aminosilicone, makes it possible to solve the technical problems mentioned above. In particular, this type of oxidizing composition improves the curl hold obtained over time, substantially reduces the porosity of permanent-waved hair and improves the compatibility of permanent-waved hair with respect to subsequent treatments.

These discoveries form the basis of the present invention.

(a) at least one enzyme of 2-electron oxidoreductase type in the presence of at least one donor for the said enzyme,

The 2-electron oxidoreductase(s) used in the oxidizing compositions in accordance with the invention can be chosen in particular from pyranose oxidases, glucose oxidases, glycerol oxidases, lactate oxidases, pyruvate oxidases and uricases.

10 By way of example, mention may be made in particular of uricase extracted from boar liver, uricase from *Arthrobacter globiformis*, as well as uricase from *Aspergillus flavus*.

The 2-electron oxidoreductase(s) in accordance with the invention preferably represent(s) from 0.01 to 20% by weight approximately relative to the total weight of the composition, and even more preferably from 0.1 to 5% by weight approximately relative to this weight.

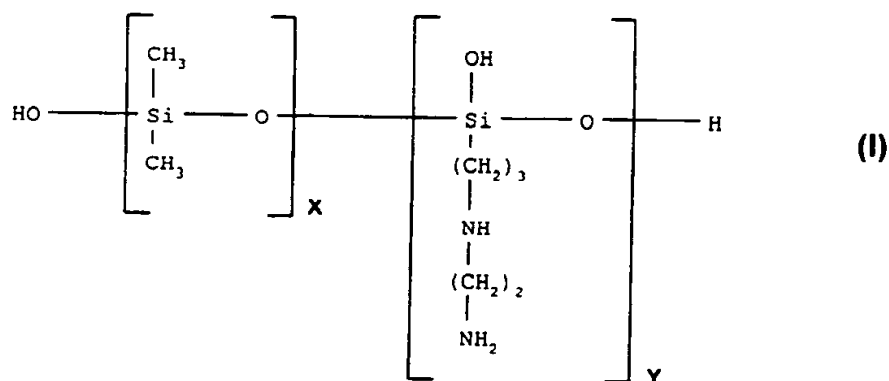
The nature of the donor (or substrate) for the said enzyme varies depending on the nature of the 2-electron oxidoreductase used. For example, as donors for the pyranose oxidases, mention may be made of D-glucose, L-sorbose and D-xylose; as a donor for the glucose oxidases, mention may be made of D-glucose; as donors for the glycerol oxidases, mention may be made of glycerol and dihydroxyacetone; as donors for the lactate oxidases, mention may be made of lactic acid and its salts; as donors for the pyruvate oxidases, mention may be made of pyruvic acid and its salts; and

lastly, as donors for the uricases, mention may be made of uric acid and its salts.

The donor(s) (or substrate(s)) used in accordance with the invention preferably represent(s) from 0.01 to 20% by weight approximately relative to the total weight of the composition in accordance with the invention, and even more preferably from 0.1 to 5% approximately relative to this weight.

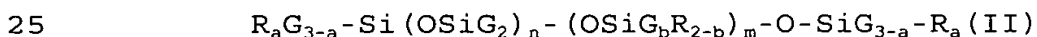
According to the invention, the term aminosilicone is understood to denote any silicone containing at least one primary, secondary or tertiary amine or a quaternary ammonium group. Mention may thus be made of:

(a) the polysiloxanes referred to in the CTFA dictionary as "amodimethicone" and corresponding to formula (I):

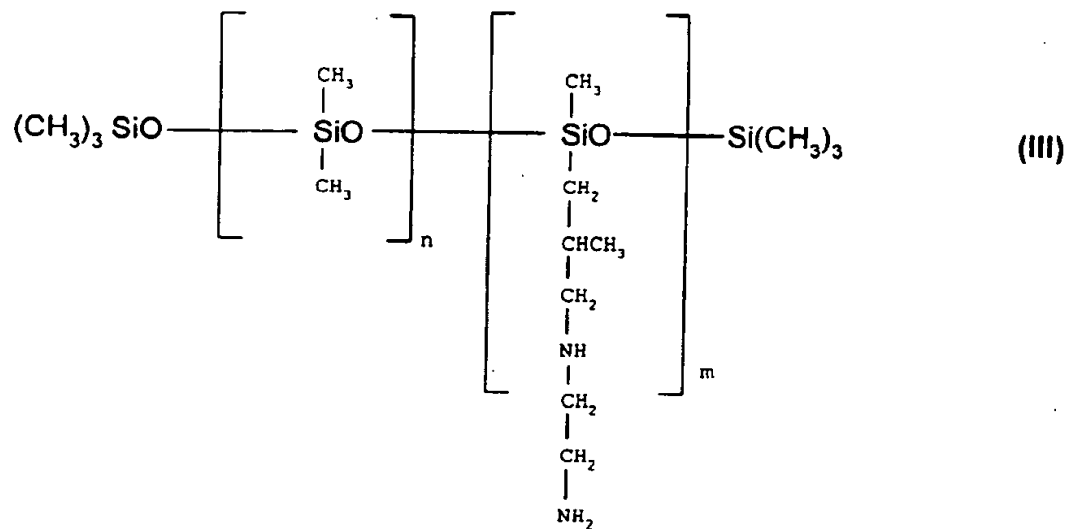


in which x and y are integers dependent on the molecular weight, generally such that the said number-average molecular weight is between 5000 and 500,000 approximately;

(b) cationic silicone polymers corresponding to the formula:

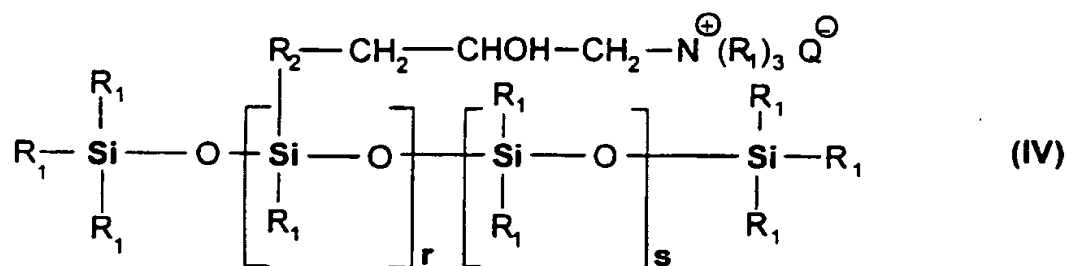


in which:



in which n and m have the meanings given above for formula (II). Such polymers are described, for example, in patent application EP-A-95238.

(c) cationic silicone polymers corresponding to the formula:



10 in which

R₁ represents a monovalent hydrocarbon-based radical containing from 1 to 18 carbon atoms, and in particular

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a C₁-C₁₈ alkyl radical or a C₂-C₁₈ alkenyl radical, for example methyl;

R₂ represents a divalent hydrocarbon-based radical, in particular a C₁-C₁₈ alkylene radical or a C₁-C₁₈, for example C₁-C₈, divalent alkyleneoxy radical;

Q⁻ is a halide ion, in particular chloride;

r represents an average statistical value from 2 to 20 and in particular from 2 to 8;

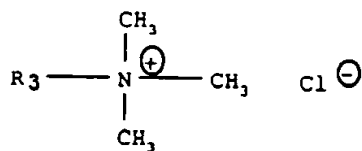
s represents an average statistical value from 20 to 200 and in particular from 20 to 50.

Such polymers are described more particularly in US patent 4,185,087.

One polymer belonging to this class is the polymer sold by the company Union Carbide under the name "Ucar Silicone ALE 56".

When these silicone polymers are used, one particularly advantageous embodiment is their use together with cationic and/or nonionic surfactants.

It is possible, for example, to use the product sold under the name "Cationic Emulsion DC 929" by the company Dow Corning, which comprises, besides amodimethicone, a cationic surfactant comprising a mixture of products corresponding to the formula:



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in which R₃ denotes alkenyl and/or alkyl radicals containing from 14 to 22 carbon atoms, derived from tallow fatty acids, and known under the CTFA name "tallowtrimonium chloride", in combination with a nonionic surfactant of formula: C₉H₁₉-C₆H₄-(OC₂H₄)₁₀-OH, known under the name "Nonoxynol 10".

Use may also be made, for example, of the product sold under the name "Cationic Emulsion DC 939" by the

company Dow Corning, which comprises, besides amodimethicone, a cationic surfactant which is trimethylcetylammmonium chloride, in combination with a nonionic surfactant of formula $C_{13}H_{27}-(OC_2H_4)_{12}-OH$, known under the CTFA name "trideceth-12".

Another commercial product which can be used according to the invention is the product sold under the name "Dow Corning Q2 7224" by the company Dow Corning, containing, in combination, trimethyl-silylamodimethicone of formula (III) described above, a nonionic surfactant of formula: $C_8H_{17}-C_6H_4-(OCH_2CH_2)_{40}-OH$, known under the CTFA name "octoxynol-40", a second nonionic surfactant of formula: $C_{12}H_{25}-(OCH_2-CH_2)_6-OH$, known under the CTFA name "isolaureth-6", and propylene glycol.

The compositions in accordance with the invention contain the aminosilicones defined above at weight contents which can be between 0.05% and 10%, preferably between 0.1% and 5% and even more preferably between 0.2% and 3%, relative to the total weight of the composition.

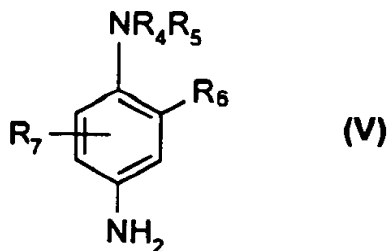
A subject of the present invention is also a ready-to-use composition for the oxidation dyeing of keratin fibres, and in particular human keratin fibres such as the hair, of the type comprising, in a medium which is suitable for dyeing, at least one oxidation base and, where appropriate, one or more couplers, which is characterized in that it contains:

- (a) at least one enzyme of 2-electron oxidoreductase type in the presence of at least one donor for the said enzyme,
- (b) at least one aminosilicone.

The nature of the oxidation base(s) used in the ready-to-use dye composition is not a critical factor. They can be chosen, in particular, from para-phenylenediamines, double bases, para-aminophenols, ortho-aminophenols and heterocyclic oxidation bases.

Among the para-phenylenediamines which can be used as oxidation bases in the dye compositions in accordance with the invention, mention may be made in

particular of the compounds of formula (V) below, and the addition salts thereof with an acid:



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in which:

- R_4 represents a hydrogen atom, a C_1 - C_4 alkyl radical, a C_1 - C_4 monohydroxyalkyl radical, a C_2 - C_4 polyhydroxyalkyl radical, a $(C_1$ - $C_4)$ alkoxy(C_1 - C_4)alkyl radical, a C_1 - C_4 alkyl radical substituted with a nitrogenous group, a phenyl radical or a 4'-aminophenyl radical;
- R_5 represents a hydrogen atom, a C_1 - C_4 alkyl radical, a C_1 - C_4 monohydroxyalkyl radical, a C_2 - C_4 polyhydroxyalkyl radical, a $(C_1$ - $C_4)$ alkoxy(C_1 - C_4)alkyl radical or a C_1 - C_4 alkyl radical substituted with a nitrogenous group;
- R_6 represents a hydrogen atom, a halogen atom such as a chlorine, bromine, iodine or fluorine atom, a C_1 - C_4 alkyl radical, a C_1 - C_4 monohydroxyalkyl radical, a C_1 - C_4 hydroxyalkoxy radical, an acetylamino(C_1 - C_4)alkoxy radical, a C_1 - C_4 mesylaminoalkoxy radical or a carbamoylamino(C_1 - C_4)alkoxy radical,
- R_7 represents a hydrogen or halogen atom or a C_1 - C_4 alkyl radical.

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Among the nitrogenous groups of formula (V) above, mention may be made in particular of amino, mono(C_1 - C_4)alkylamino, di(C_1 - C_4)alkylamino, tri(C_1 - C_4)alkylamino, monohydroxy(C_1 - C_4)alkylamino, imidazolinium and ammonium radicals.

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Among the para-phenylenediamines of formula (V)

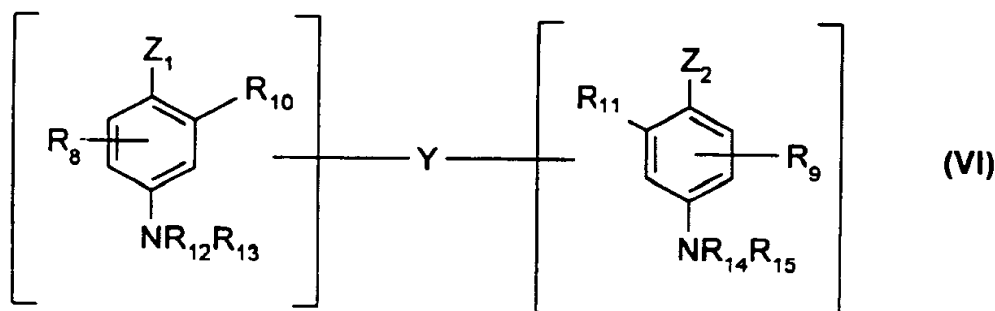
above, mention may be made more particularly of para-phenylenediamine, para-toluylenediamine, 2-chloro-para-phenylenediamine, 2,3-dimethyl-para-phenylenediamine, 2,6-dimethyl-para-phenylenediamine, 2,6-diethyl-para-phenylenediamine, 2,5-dimethyl-para-phenylenediamine, N,N-dimethyl-para-phenylenediamine, N,N-diethyl-para-phenylenediamine, N,N-dipropyl-para-phenylenediamine, 4-amino-N,N-diethyl-3-methylaniline, N,N-bis(β -hydroxyethyl)-para-phenylenediamine, 4-amino-N,N-bis(β -hydroxyethyl)-2-methylaniline, 4-amino-2-chloro-N,N-bis(β -hydroxyethyl)aniline, 2- β -hydroxyethyl-para-phenylenediamine, 2-fluoro-para-phenylenediamine, 2-isopropyl-para-phenylenediamine, N-(β -hydroxypropyl)-para-phenylenediamine, 2-hydroxymethyl-para-phenylenediamine, N,N-dimethyl-3-methyl-para-phenylenediamine, N,N-(ethyl- β -hydroxyethyl)-para-phenylenediamine, N-(β,γ -dihydroxypropyl)-para-phenylenediamine, N-(4'-aminophenyl)-para-phenylenediamine, N-phenyl-para-phenylenediamine, 2- β -hydroxyethyloxy-para-phenylenediamine, 2- β -acetylaminoethyloxy-para-phenylenediamine and N-(β -methoxyethyl)-para-phenylenediamine, and the addition salts thereof with an acid.

Among the para-phenylenediamines of formula (V) above, para-phenylenediamine, para-toluylenediamine, 2-isopropyl-para-phenylenediamine, 2- β -hydroxyethyl-para-phenylenediamine, 2- β -hydroxyethyloxy-para-phenylenediamine, 2,6-dimethyl-para-phenylenediamine, 2,6-diethyl-para-phenylenediamine, 2,3-dimethyl-para-phenylenediamine, N,N-bis(β -hydroxyethyl)-para-phenylenediamine, 2-chloro-para-phenylenediamine and 2- β -acetylaminoethyloxy-para-phenylenediamine and the addition salts thereof with an acid are most particularly preferred.

According to the invention, the term double bases is understood to refer to the compounds containing at least two aromatic rings bearing amino and/or hydroxyl groups.

Among the double bases which can be used as oxidation bases in the dye compositions in accordance with the invention, mention may be made in particular of the compounds corresponding to formula (VI) below, and the addition salts thereof with an acid:

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in which:

- 15 - Z_1 and Z_2 , which may be identical or different, represent a hydroxyl or $-NH_2$ radical which may be substituted with a C_1-C_4 alkyl radical or with a linker arm Y;
- the linker arm Y represents a linear or branched
- 20 alkylene chain containing from 1 to 14 carbon atoms, which may be interrupted by or terminated with one or more nitrogenous groups and/or one or more hetero atoms such as oxygen, sulphur or nitrogen atoms, and optionally substituted with one or more hydroxyl or C_1-C_6 alkoxy
- 25 radicals;
- R_8 and R_9 represent a hydrogen or halogen atom, a C_1-C_4 alkyl radical, a C_1-C_4 monohydroxyalkyl radical, a C_2-C_4

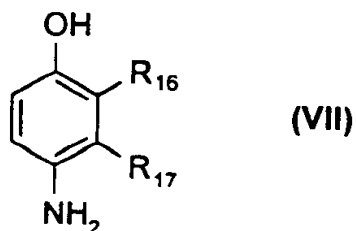
- R₁₀, R₁₁, R₁₂, R₁₃, R₁₄ and R₁₅, which may be identical or different, represent a hydrogen atom, a linker arm Y or a C₁-C₄ alkyl radical;

Among the nitrogenous groups of formula (VI) above, mention may be made in particular of amino, mono(C₁-C₄)alkylamino, di(C₁-C₄)alkylamino, tri(C₁-C₄)alkylamino, monohydroxy(C₁-C₄)alkylamino, imidazolinium and ammonium radicals.

Among the double bases of formula (VI) above, mention may be made more particularly of N,N'-bis(β -hydroxyethyl)-N,N'-bis(4'-aminophenyl)-1,3-diaminopropanol, N,N'-bis(β -hydroxyethyl)-N,N'-bis(4'-aminophenyl)ethylenediamine, N,N'-bis(4-aminophenyl)-tetramethylenediamine, N,N'-bis(β -hydroxyethyl)-N,N'-bis(4-aminophenyl)tetramethylenediamine, N,N'-bis(4-methylaminophenyl)tetramethylenediamine, N,N'-bis(ethyl)-N,N'-bis(4'-amino-3'-methylphenyl)ethylenediamine and 1,8-bis(2,5-diaminophenoxy)-3,5-dioxaoctane, and the addition salts thereof with an acid.

Among these double bases of formula (VI), N,N'-bis(β -hydroxyethyl)-N,N'-bis(4'-aminophenyl)-1,3-diaminopropanol and 1,8-bis(2,5-diaminophenoxy)-3,5-dioxaoctane, or one of the addition salts thereof with an acid, are particularly preferred.

Among the para-aminophenols which can be used as
30 oxidation bases in the dye compositions in accordance
with the invention, mention may be made in particular of
the compounds corresponding to formula (VII) below, and
the addition salts thereof with an acid:



in which:

- R_{16} represents a hydrogen or halogen atom or a C_1 - C_4 alkyl, C_1 - C_4 monohydroxyalkyl, $(C_1$ - C_4)alkoxy(C_1 - C_4)alkyl, C_1 - C_4 aminoalkyl or hydroxy(C_1 - C_4)alkylamino- (C_1 - C_4)alkyl radical,

- R_{17} represents a hydrogen or halogen atom or a C_1 - C_4 -alkyl, C_1 - C_4 monohydroxyalkyl, C_2 - C_4 polyhydroxyalkyl, C_1 - C_4 aminoalkyl, C_1 - C_4 cyanoalkyl or $(C_1$ - C_4)alkoxy- (C_1 - C_4)alkyl radical,

it being understood that at least one of the radicals R_{16} or R_{17} represents a hydrogen atom.

Among the para-aminophenols of formula (VII) above, mention may be made more particularly of para-aminophenol, 4-amino-3-methylphenol, 4-amino-3-fluorophenol, 4-amino-3-hydroxymethylphenol, 4-amino-2-methylphenol, 4-amino-2-hydroxymethylphenol, 4-amino-2-methoxymethylphenol, 4-amino-2-aminomethylphenol, 4-amino-2-(β -hydroxyethylaminomethyl)phenol and 4-amino-2-fluorophenol, and the addition salts thereof with an acid.

Among the ortho-aminophenols which can be used as oxidation bases in the dye compositions in accordance with the invention, mention may be made more particularly of 2-aminophenol, 2-amino-5-methylphenol, 2-amino-6-methylphenol and 5-acetamido-2-aminophenol, and the addition salts thereof with an acid.

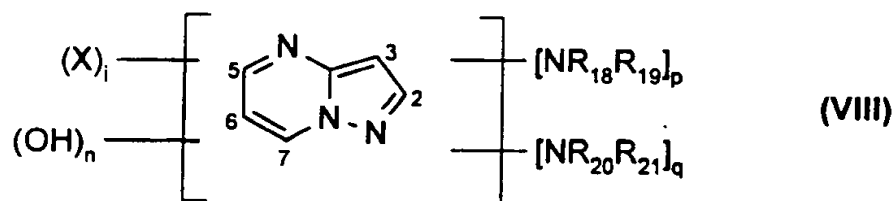
Among the heterocyclic bases which can be used as oxidation bases in the dye compositions in accordance with the invention, mention may be made more particularly

Among the pyridine derivatives, mention may be made more particularly of the compounds described, for example, in patents GB 1,026,978 and GB 1,153,196, such as 2,5-diaminopyridine, 2-(4-methoxyphenyl)amino-3-aminopyridine, 2,3-diamino-6-methoxypyridine, 2-(β -methoxyethyl)amino-3-amino-6-methoxypyridine and 3,4-diaminopyridine, and the addition salts thereof with an acid.

20 Among the pyrazole derivatives, mention may be
made more particularly of the compounds described in
patents DE 3,843,892, DE 4,133,957 and patent
applications WO 94/08969, WO 94/08970, FR-A-2,733,749 and
DE 195 43 988, such as 4,5-diamino-1-methylpyrazole, 3,4-
25 diaminopyrazole, 4,5-diamino-1-(4'-chlorobenzyl)pyrazole,
4,5-diamino-1,3-dimethylpyrazole, 4,5-diamino-3-methyl-1-
phenylpyrazole, 4,5-diamino-1-methyl-3-phenylpyrazole, 4-
amino-1,3-dimethyl-5-hydrazinopyrazole, 1-benzyl-4,5-
diamino-3-methylpyrazole, 4,5-diamino-3-tert-butyl-1-
30 methylpyrazole, 4,5-diamino-1-tert-butyl-3-
methylpyrazole, 4,5-diamino-1-(β -hydroxyethyl)-3-
methylpyrazole, 4,5-diamino-1-ethyl-3-methylpyrazole,
4,5-diamino-1-ethyl-3-(4'-methoxyphenyl)pyrazole, 4,5-
diamino-1-ethyl-3-hydroxymethylpyrazole, 4,5-diamino-3-
35 hydroxymethyl-1-methylpyrazole, 4,5-diamino-3-
hydroxymethyl-1-isopropylpyrazole, 4,5-diamino-3-methyl-
1-isopropylpyrazole, 4-amino-5-(2'-aminoethyl)amino-1,3-
dimethylpyrazole, 3,4,5-triaminopyrazole, 1-methyl-3,4,5-

triaminopyrazole, 3,5-diamino-1-methyl-4-methylamino-pyrazole and 3,5-diamino-4-(β -hydroxyethyl)amino-1-methylpyrazole, and the addition salts thereof with an acid.

5 Among the pyrazolopyrimidine derivatives, mention may be made more particularly of the pyrazolo[1,5-a]pyrimidines of formula (VIII) below, and the addition salts thereof with an acid or with a base and the tautomeric forms thereof, when a tautomeric equilibrium
10 exists:

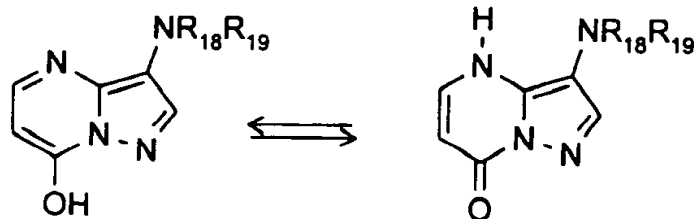


in which:

- 15 - R_{18} , R_{19} , R_{20} and R_{21} , which may be identical or different, denote a hydrogen atom, a C_1 - C_4 alkyl radical, an aryl radical, a C_1 - C_4 hydroxyalkyl radical, a C_2 - C_4 polyhydroxyalkyl radical, a $(C_1$ - C_4)alkoxy(C_1 - C_4)alkyl radical, a C_1 - C_4 aminoalkyl radical (it being possible for
20 the amine to be protected with an acetyl, ureido or sulphonyl radical), a $(C_1$ - C_4)alkylamino(C_1 - C_4)alkyl radical, a di[(C_1 - C_4)alkyl]amino(C_1 - C_4)alkyl radical (it being possible for the dialkyl radicals to form a 5- or 6-membered carbon-based ring or a heterocycle), a
25 hydroxy(C_1 - C_4)alkyl- or di[hydroxy-(C_1 - C_4)alkyl]amino(C_1 - C_4)alkyl radical;
- the radicals X, which may be identical or different, denote a hydrogen atom, a C_1 - C_4 alkyl radical, an aryl radical, a C_1 - C_4 hydroxyalkyl radical, a C_2 - C_4 polyhydroxyalkyl radical, a C_1 - C_4 aminoalkyl radical, a
30 $(C_1$ - C_4)alkylamino(C_1 - C_4)alkyl radical, a di[(C_1 - C_4)alkyl]amino(C_1 - C_4)alkyl radical (it being possible for

- the dialkyls to form a 5- or 6-membered carbon-based ring or a heterocycle), a hydroxy(C₁-C₄)alkyl- or di[hydroxy(C₁-C₄)alkyl]amino(C₁-C₄)alkyl radical, an amino radical, a (C₁-C₄)alkyl- or di[(C₁-C₄)alkyl]amino radical; a halogen atom, a carboxylic acid group or a sulphonic acid group;
- i is equal to 0, 1, 2 or 3;
 - p is equal to 0 or 1;
 - q is equal to 0 or 1;
 - n is equal to 0 or 1;
- with the proviso that:
- the sum p + q is other than 0;
 - when p + q is equal to 2, then n is equal to 0 and the groups NR₁₈R₁₉ and NR₂₀R₂₁ occupy the (2,3); (5,6); (6,7); (3,5) or (3,7) positions;
 - when p + q is equal to 1, then n is equal to 1 and the group NR₁₈R₁₉ (or NR₂₀R₂₁) and the OH group occupy the (2,3); (5,6); (6,7); (3,5) or (3,7) positions.

When the pyrazolo[1,5-a]pyrimidines of formula (VIII) above are such that they contain a hydroxyl group on one of the positions 2, 5 or 7 α to a nitrogen atom, a tautomeric equilibrium exists represented, for example, by the following scheme:



- Among the pyrazolo[1,5-a]pyrimidines of formula (VIII) above, mention may be made in particular of:
- pyrazolo[1,5-a]pyrimidine-3,7-diamine;
 - 2,5-dimethylpyrazolo[1,5-a]pyrimidine-3,7-diamine;
 - pyrazolo[1,5-a]pyrimidine-3,5-diamine;
 - 2,7-dimethylpyrazolo[1,5-a]pyrimidine-3,5-diamine;

- 3-aminopyrazolo[1,5-a]pyrimidin-7-ol;
- 3-aminopyrazolo[1,5-a]pyrimidin-5-ol;
- 2-(3-aminopyrazolo[1,5-a]pyrimidin-7-ylamino)ethanol;
- 2-(7-aminopyrazolo[1,5-a]pyrimidin-3-ylamino)ethanol;
- 5 - 2-[(3-aminopyrazolo[1,5-a]pyrimidin-7-yl)-(2-hydroxyethyl)amino]ethanol;
- 2-[(7-aminopyrazolo[1,5-a]pyrimidin-3-yl)-(2-hydroxyethyl)amino]ethanol;
- 5,6-dimethylpyrazolo[1,5-a]pyrimidine-3,7-diamine;
- 10 - 2,6-dimethylpyrazolo[1,5-a]pyrimidine-3,7-diamine;
- 2,5,N7,N7-tetramethylpyrazolo[1,5-a]pyrimidine-3,7-diamine;

and the addition salts thereof and the tautomeric forms thereof, when a tautomeric equilibrium exists.

- 15 The pyrazolo[1,5-a]pyrimidines of formula (VIII) above can be prepared by cyclization starting with an aminopyrazole, according to the syntheses described in the following references:

- EP 628559 Beiersdorf-Lilly.
- 20 - R. Vishdu, H. Navedul, Indian J. Chem., 34b (6), 514, 1995.
- N.S. Ibrahim, K.U. Sadek, F.A. Abdel-Al, Arch. Pharm., 320, 240, 1987.
- R.H. Springer, M.B. Scholten, D.E. O'Brien,
- 25 T. Novinson, J.P. Miller, R.K. Robins, J. Med. Chem., 25, 235, 1982.
- T. Novinson, R.K. Robins, T.R. Matthews, J. Med. Chem., 20, 296, 1977.
- US 3907799 ICN Pharmaceuticals.

- 30 The pyrazolo[1,5-a]pyrimidines of formula (VIII) above can also be prepared by cyclization starting from hydrazine, according to the syntheses described in the following references:

- A. McKillop and R.J. Kobilecki, Heterocycles, 6(9),
- 35 1355, 1977.
- E. Alcade, J. De Mendoza, J.M. Marcia-Marquina, C. Almera, J. Elguero, J. Heterocyclic Chem., 11(3), 423, 1974.

- K. Saito, I. Hori, M. Higashashi, H. Midorikawa, Bull. Chem. Soc. Japan, 47(2), 476, 1974.

The oxidation base(s) in accordance with the invention preferably represent(s) from 0.0005 to 12% by weight approximately relative to the total weight of the ready-to-use dye composition, and even more preferably from 0.005 to 6% by weight approximately relative to this weight.

The couplers which can be used are those used conventionally in oxidation dye compositions, i.e. meta-phenylenediamines, meta-aminophenols and meta-diphenols, mono- or polyhydroxylated naphthalene derivatives, sesamol and its derivatives and heterocyclic compounds such as, for example, indole derivatives, indoline derivatives, benzimidazole derivatives, benzomorpholine derivatives, sesamol derivatives, pyrazoloazole derivatives, pyrroloazole derivatives, imidazoloazole derivatives, pyrazolopyrimidine derivatives, pyrazoline-3,5-dione derivatives, pyrrolo[3,2-d]oxazole derivatives, pyrazolo[3,4-d]thiazole derivatives, thiazoloazole S-oxide derivatives and thiazoloazole S,S-dioxide derivatives, and the addition salts thereof with an acid.

These couplers can be chosen in particular from 2-methyl-5-aminophenol, 5-N-(β -hydroxyethyl)amino-2-methylphenol, 3-aminophenol, 1,3-dihydroxybenzene, 1,3-dihydroxy-2-methylbenzene, 4-chloro-1,3-dihydroxybenzene, 2,4-diamino-1-(β -hydroxyethyloxy)benzene, 2-amino-4-(β -hydroxyethylamino)-1-methoxybenzene, 1,3-diaminobenzene, 1,3-bis(2,4-diaminophenoxy)propane, sesamol, α -naphthol, 6-hydroxyindole, 4-hydroxyindole, 4-hydroxy-N-methylindole, 6-hydroxyindoline, 2,6-dihydroxy-4-methylpyridine, 1H-3-methylpyrazol-5-one and 1-phenyl-3-methylpyrazol-5-one, and the addition salts thereof with an acid.

35 When they are present, these couplers preferably
represent from 0.0001 to 10% by weight approximately
relative to the total weight of the ready-to-use dye

composition, and even more preferably from 0.005 to 5% by weight approximately relative to this weight.

In general, the addition salts with an acid which can be used in the context of the dye compositions of the invention (oxidation bases and couplers) are chosen in particular from the hydrochlorides, hydrobromides, sulphates, tartrates, lactates and acetates.

The dye composition of the invention can also contain, in addition to the oxidation dye precursors defined above and the optional combined couplers, direct dyes to enrich the shades with glints. These direct dyes can then be chosen in particular from nitro dyes, azo dyes or anthraquinone dyes.

The subject of the invention is also a process for dyeing keratin fibres, and in particular human keratin fibres such as the hair, using the ready-to-use dye composition as defined above.

According to this process, at least one ready-to-use dye composition as defined above is applied to the fibres, for a period which is sufficient to develop the desired coloration, after which the fibres are rinsed, optionally washed with shampoo, rinsed again and dried.

The time required to develop the coloration on the keratin fibres is generally between 3 and 60 minutes and even more precisely between 5 and 40 minutes.

According to one specific embodiment of the invention, the process includes a first step which consists in separately storing, on the one hand, a composition (A) comprising, in a medium which is suitable for dyeing, at least one oxidation base and optionally at least one coupler as defined above, and, on the other hand, a composition (B) containing, in a medium which is suitable for dyeing, at least one enzyme of 2-electron oxidoreductase type in the presence of at least one donor for the said enzyme and at least one aminosilicone, and then in mixing them together at the time of use, before applying this mixture to the keratin fibres.

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Another subject of the invention is a multi-compartment dyeing device or "kit" or any other multi-compartment packaging system, a first compartment of which contains composition (A) as defined above and a second compartment of which contains composition (B) as defined above. These devices can be equipped with means for applying the desired mixture to the hair, such as the devices described in patent FR-2,586,913 in the name of the Applicant.

The first step (i) of this process consists in applying a reducing composition to the hair. This application is carried out lock by lock or all at once.

The usual step for placing the hair under tension
35 in a shape corresponding to the desired final shape for
this hair (for example curls) can be carried out by any
suitable means, in particular mechanical means, which are
suitable and known per se for maintaining the hair under

The hair can also be shaped without the aid of external means, simply with the fingers.

In the optional second step of the process (step
15 (ii)), the hair impregnated with the reducing composition
is then rinsed thoroughly with an aqueous composition.

As in the case of the application of the reducing composition, the hair onto which the oxidizing composition has been applied is then, conventionally, left for a standing or waiting phase lasting a few minutes, generally between 3 and 30 minutes, preferably between 5 and 15 minutes.

Lastly, in the final step of the process according to the invention (step (iv)), which is also optional, the hair impregnated with the oxidizing composition is rinsed thoroughly, generally with water.

The oxidizing composition according to the invention can also be used in a process for bleaching keratin fibres, and in particular the hair.

5 Conventionally, a second step of the bleaching process according to the invention is a step of rinsing the keratin fibres.

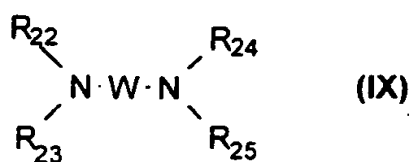
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The pH of the ready-to-use dye compositions and of the oxidizing compositions used for the permanent reshaping or bleaching of the keratin fibres in accordance with the invention is chosen such that the enzymatic activity of the 2-electron oxidoreductase is not adversely affected. It is generally between 5 and 11 approximately, and preferably between 6.5 and 10 approximately. It can be adjusted to the desired value using acidifying or basifying agents usually used for dyeing keratin fibres.

Among the acidifying agents, mention may be made, by way of example, of inorganic or organic acids such as

hydrochloric acid, orthophosphoric acid, sulphuric acid, carboxylic acids such as acetic acid, tartaric acid, citric acid or lactic acid, and sulphonc acids.

Among the basifying agents, mention may be made, by way of example, of aqueous ammonia, alkaline carbonates, alkanolamines such as mono-, di- and triethanolamines, 2-methyl-2-aminopropanol and derivatives thereof, sodium hydroxide, potassium hydroxide and the compounds of formula (IX) below:



in which W is a propylene residue optionally substituted with a hydroxyl group or a C₁-C₄ alkyl radical; R₂₂, R₂₃, R₂₄ and R₂₅, which may be identical or different, represent a hydrogen atom or a C₁-C₄ alkyl or C₁-C₄ hydroxyalkyl radical.

The ready-to-use dye compositions and the oxidizing compositions for the permanent reshaping or bleaching of keratin fibres in accordance with the invention can also contain various adjuvants used conventionally in compositions for dyeing, permanently reshaping or bleaching the hair, such as anionic, cationic, nonionic, amphoteric or zwitterionic surfactants or mixtures thereof, anionic, cationic, nonionic, amphoteric or zwitterionic polymers or mixtures thereof, inorganic or organic thickeners, antioxidants, enzymes other than the 2-electron oxidoreductases used in accordance with the invention, such as, for example, peroxidases, penetration agents, sequestering agents, fragrances, buffers, dispersing agents, conditioners, film-forming agents, preserving agents and opacifiers.

Needless to say, a person skilled in the art will take care to select this or these optional complementary

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on a weight basis.

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Example 1 of dye composition:

The ready-to-use dye composition below was prepared (contents in grams):

- 5
- Uricase from *Arthrobacter globiformis* at a concentration of 20 International Units (I.U.)/mg, sold by the company Sigma 1.5 g
 - Uric acid 1.5 g
 - para-Phenylenediamine 0.324 g
 - Resorcinol 0.33 g
 - 10 - Polydimethylsiloxane containing amino-ethylaminopropyl groups, at a concentration of 35% in water, sold under the name Dow Corning 939 Emulsion 1.2 g
 - Demineralized water qs 100 g

15 The ready-to-use dye composition described above was applied to locks of natural grey hair containing 90% white hairs for 30 minutes. The hair was then rinsed, washed with a standard shampoo and then dried.

20 Locks of hair dyed a matt dark-blonde colour were obtained.

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Example 2: Oxidizing composition for permanent-waving
or bleaching

- | | | |
|----|--|-----------|
| | - Uricase from <i>Arthrobacter globiformis</i> at a concentration of 20 International Units (I.U.)/mg, sold by the company Sigma | 1.8 g |
| 5 | - Uric acid | 1.65 g |
| | - Ethanol | 20.0 g |
| | - (C ₈ -C ₁₀)alkyl polyglucoside as an aqueous solution containing 60% active material (A.M.), sold under the name Oramix CG110 by the company SEPPIC | 8.0 g |
| | - Mixture of polydimethylsiloxane containing aminoethylaminoisobutyl/polydimethylsiloxane groups, sold under the trade name Dow Corning Q2 8220 Fluid | 1.0 g |
| | - 2-Methy-2-amino-1-propanol | qs pH 9.5 |
| 10 | - Demineralized water | qs 100 g |